SEMICONDUCTOR DEVICE AND MANUFACTURING METHOD OF THE SAME

CROSS REFERENCE TO RELATED APPLICATIONS

5   This application is based upon and claims the benefit of priority from prior Japanese Patent Application P2002-278088 filed on September 24, 2002; the entire contents of which are incorporated by reference herein.

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BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates to a semiconductor device and a manufacturing method for the same. In particular, it relates

15   to a semiconductor device using a semiconductor layer doped with impurity atoms having different covalent bond radii.

2. Description of the Related Art

Control of an impurity distribution in a semiconductor layer

20   is extremely important for a semiconductor device. Generally, a thermal diffusion process is used as a technique to form an impurity distribution in a semiconductor layer with high controllability. However, depending on a doping impurity atom, an insufficient concentration of an electrically active impurity atom is provided.

25   Alternatively, the desired impurity distribution is not provided which results in a large diffusion coefficient of the impurity

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atom. For example, in order to suppress a short channel effect and achieve a high drive current in a metal-oxide-semiconductor field effect transistor (MOSFET) fabricated on a semiconductor substrate such as silicon (Si), it is necessary to form a super

5   steep retrograde channel profile (SSRP), where impurity concentration decreases rapidly toward a gate oxide layer in a channel region of the MOSFET. Otherwise, in a source/drain region, it is required to form a shallow impurity diffusion layer with low resistance.

10   Indium (In) is used as an acceptor impurity atom in order to form the SSRP of an n-channel MOSFET. However, it is difficult to form a high carrier concentration in a deep channel region because of a low active impurity concentration of In. In addition, in a source/drain region of a p-channel MOSFET, boron (B) is used

15   as an acceptor impurity atom. However, it is very difficult to form a shallow impurity diffusion layer so that a diffusion coefficient of B in a Si crystal is large. Further, because of the low active impurity concentration, In cannot be used as an acceptor impurity atom of the source/drain region of the p-channel

20   MOSFET.

Many proposals for a technique such as "co-doping" which dopes plural impurity atoms together in a semiconductor layer, have been disclosed. According to the co-doping technique, in order to reduce the crystal defect in a semiconductor layer by

25   a vapor phase growth, a method in which phosphorus (P) and arsenic (As) are doped together in a Si layer has been reported (see Japanese

Patent Application Nos. 55-028215 and 55-025492). However, a carrier concentration cannot exceed the active impurity concentration of P and As. In addition, in order to suppress diffusion of an acceptor atom, a method to co-dope a constituent

5   semiconductor atom in addition to the acceptor atom has been reported (see Japanese Patent Laid-Open No. 2000-68225). In addition, when doping B into the Si layer, a method to co-dope germanium (Ge) is used. However, it is necessary to dope Ge with a high concentration to change the activation rate and the diffusion for

10   B.

As described above, in a semiconductor device such as the MOSFET, the doping layer of the high impurity concentration is needed in the channel or the source/drain region. However, it is impossible to achieve a higher carrier concentration exceeding

15   the active impurity concentration of the dopant atom. It is also impossible to suppress the diffusion of the heavily doped impurity atom.

20   SUMMARY OF THE INVENTION

An object of the present invention is to provide a semiconductor device having a doping layer so as to improve an activation rate of an impurity atom and to suppress diffusion of the impurity atom.

25   A first aspect of the present invention inheres in a semiconductor device having a semiconductor layer including a first

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impurity atom having a covalent bond radius larger than a minimum radius of a covalent bond of a semiconductor constituent atom of a semiconductor layer, and a second impurity atom having a covalent bond radius smaller than a maximum radius of the covalent

5 bond of the semiconductor constituent atom, wherein the first and second impurity atoms are arranged in a nearest neighbor lattice site location and at least one of the first and second impurity atoms is electrically active.

A second aspect of the present invention inheres in a

10 manufacturing method of a semiconductor device including providing a semiconductor substrate, doping a first impurity atom having a covalent bond radius larger than a minimum radius of a covalent bond of a semiconductor constituent atom of a semiconductor layer of the semiconductor substrate, and doping a second impurity atom

15 having a covalent bond radius smaller than a maximum radius of a covalent bond of the semiconductor constituent atom so as to be arranged in a nearest neighbor lattice site of the first impurity atom.

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BRIEF DESCRIPTION OF DRAWINGS

Fig. 1 is a cross-sectional view showing an example of a semiconductor layer of a semiconductor device according to an embodiment of the present invention;

25 Fig. 2 is a figure showing an electronic state of a semiconductor by the tight-binding approximation description according to an

embodiment of the present invention;

Fig. 3 is a figure showing bonding states of electronic orbital of Si by the tight-binding approximation description according to the embodiment of the present invention;

5   Figs. 4A and 4B are schematic diagrams explaining bonding states of nearest neighbor impurity atoms in a semiconductor according to the embodiment of the present invention;

Figs. 5A and 5B are schematic diagrams showing energy levels of bonding states of a cluster between the same impurity atoms

10   in a semiconductor according to the embodiment of the present invention;

Figs. 6A and 6B are schematic diagrams showing energy levels of bonding states of a cluster between the first and second impurity atoms in a semiconductor according to the embodiment of the present

15   invention;

Fig. 7 is a figure showing an example of impurity distributions in a semiconductor layer according to an embodiment of the present invention; and

Figs. 8A through 8F are process diagrams illustrating a

20   manufacturing method of a semiconductor device according to the embodiment of the present invention.

DETAILED DESCRIPTION OF THE INVENTION

25   Various embodiments of the present invention will be described with reference to the accompanying drawings. It is to

be noted that the same or similar reference numerals are applied to the same or similar parts and elements throughout the drawings, and the description of the same or similar parts and elements will be omitted or simplified.

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A semiconductor device according to an embodiment of the present invention includes, as shown in Fig. 1, a semiconductor layer 12 doped with first and second impurity atoms by a nearest neighbor lattice site location in a principal surface side of

10   a semiconductor substrate 10. On the semiconductor layer 12, for example, an insulating film 14 such as a gate oxide film of the semiconductor device is deposited. For example, a covalent bond radius of the first impurity atom is larger than the minimum of a covalent bond radius of a constituent atom of

15   the semiconductor layer 12, and a covalent bond radius of the second impurity atom is smaller than the maximum of the covalent bond radius of the constituent atom of the semiconductor layer 12. At least one of the first and second impurity atoms is an acceptor or a donor impurity atom.

20   At first a base model of the embodiment of the present invention is explained by using Fig. 2 through Fig. 6. By simulating an electron bonding orbital using the tight-binding approximation in a semiconductor having a tetrahedral arrangement such as diamond (C), silicon (Si), germanium (Ge),

25   a column III-V compound semiconductor and a column II-VI compound semiconductor, a semiconductor atom 51 arranges an sp^3 hybridized

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orbital 53, as shown in Fig. 2. When the semiconductor atom 51 assumes a molecule state from an independent atomic state, as shown in Fig. 3, the sp^3 hybridized orbital 53 of a orbital energy level Eh of the independent atomic state is separated

5 into a bonding orbital of a bonding energy level E_B between adjacent semiconductor atoms 51 and an anti-bonding orbital of an anti-bonding energy level E_{AB}. Furthermore, when the molecule state is changed to a crystal state, the bonding orbital and the anti-bonding orbital are expanded to form a valence band

10 VB and a conduction band CB having a band gap energy Eg.

An electron configuration where two impurity atoms serving as the acceptor or the donor, producing a comparatively deep impurity level in the gap are arranged in a nearest neighbor substitutional site is considered. As an operative example to

15 simplify explanation, In and B which are acceptors in a column IV element semiconductor such as Si are described. However, other acceptor impurity atoms may be used. In addition, similar considerations are appropriate in the case of a donor by replacing an electron with a hole. Furthermore, similar considerations

20 are appropriate for other semiconductors including a compound semiconductor.

As shown in Figs. 4A and 4B, when a first impurity atom 61 and a second impurity atom 62 are arranged in a nearest neighbor substitutional site, two configurations can be considered. One

25 is, as shown in Fig. 4A, a configuration of a bonding orbital 63 between the first impurity atoms 61 and the second impurity

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atoms 62 which is like the sp^3 hybridized orbital, as well as bonding orbitals 65a or 65b between the semiconductor atom 51 and the first impurity atom 61 or the second impurity atom 62. The configuration shown in Fig. 4A is defined as "an sp^3

5   configuration". The other is, as shown in Fig. 4B, a configuration where bonding between the first impurity atom 61 and the second impurity atom 62 is broken so as to separate the impurity atoms from each other and to move respectively toward the semiconductor atom 51. The bonding orbitals 69a or 69b

10   between the semiconductor atom 51 and the first impurity atom 61 or the second impurity atom 62, are like a planar sp^2 hybridized orbital as similar to graphite. The broken bonding orbitals 67a, 67b of the first and second impurity atoms 61, 62 serve as p orbitals, respectively. The configuration shown in Fig.

15   4 (b) is defined as "an sp^2 configuration".

To begin with, the case where the first and the second impurity atoms 61, 62 are the same impurity atom, In, is described by use of Figs. 5A and 5B. Energy levels of the bonding orbitals of In-Si or In-In located between the valence band VB and the

20   conduction band CB, which are separated by the band gap energy Eg, are determined by deduction. Fig. 5A shows the energy level of the bonding orbitals when In provides the sp^3 configuration. All four In sp^3 hybridized orbitals have the same energy level EI_{sp3}. The In-Si bonding orbitals provided by the hybridized

25   orbitals with three Si atoms adjacent to the In atom, form the acceptor level E_{A1} at a comparatively shallow position near the

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upper end of the valence band VB, as is the case where the In atom is alone in a substitutional site. On the other hand, the energy level EI_{sp3} of the In sp^3 hybridized orbital is higher than the energy level ES_{sp3} of the Si sp^3 hybridized orbital.

5   Therefore, the energy level of the In-In bonding orbital for the nearest neighbor substitutional site is slightly higher than the acceptor level E_{A1} of the In-Si bonding orbital, and forms a deep acceptor level E_{D1}. In addition, because the In atom is tervalent, an electron does not provide for the deep

10   acceptor level E_{D1}.

Fig. 5B shows the energy level of the bonding orbital when In provides the sp^2 configuration. The In orbitals are, as shown in Fig. 4B, of three planar sp^2 hybridized orbitals spreading in three directions and the p orbital. In comparison with the

15   energy level EI_{sp3} of the In sp^3 hybridized orbital, the energy level EI_{sp2} of the sp^2 hybridized orbital is low, and the energy level EI_p of the p orbital is high. Therefore, the acceptor level E_{A2} of the In-Si bonding orbital of the sp^2 configuration is lower than the acceptor level E_{A1} of the sp^3 configuration.

20   On the contrary, the acceptor level E_{D2} of the p orbital of the sp^2 configuration is higher than the deep acceptor level E_{D1} of the In-In bonding orbital. Then the acceptor level E_{D2} is formed at a position near conduction band CB from the center of the band gap. Since the p orbital is empty, i.e., the In-In bond

25   is broken, electronic orbital energy of the sp^2 configuration completely decreases, as compared with the sp^3 configuration.

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The decreased energy difference is defined as an energy gain $\Delta\varepsilon$ by the sp^2 configuring.

As described above, the electronic energy of the bonding orbital near the upper end of the valence band VB decreases by shifting from the sp^3 configuration to the sp^2 configuration. Practically, an elastic potential energy increases with increment ΔEs, since the lattice is distorted by shifting from the sp3 configuration to the sp^2 configuration. Depending on the magnitude relation between the energy gain $\Delta\varepsilon$ by sp^2 configuring and the elastic potential energy increment ΔEs, the bonding orbital under consideration is provided either the sp^3 configuration or the sp^2 configuration. When the acceptor level E_{A1} is deeper, the energy gain $\Delta\varepsilon$ by sp^2 configuring increases and the sp^2 configuration easily tends to be formed. A great difference occurs in a function as a dopant in the sp^2 configuration and the sp^3 configuration. In other words, in the sp^3 configuration, the In-In bonding orbital is basically electrically active although energy increases. On the other hand, in the sp^2 configuration, the bonding orbital is electrically inactive because of the large energy increase of the bonding orbital.

The In atom has a large covalent bond radius in comparison with Si. Generally, when the impurity atom that has a different covalent bond radius from Si comes close to the same impurity atom, the elastic potential energy increases due to the lattice distortion. Therefore, total energy of the bonding orbital

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increases. However, when the energy gain $\Delta \varepsilon$ by sp^2 configuring

is large enough, the elastic potential energy ΔEs is overcome

and an In-In cluster is formed. In addition, when the acceptor

level of the lone impurity atom is deep, energy gain increases

5   by filling two electrons in the bonding orbital between the

acceptor levels of the impurity atom. Therefore, an attraction

force between the impurity atoms has an effect that the impurity

cluster become stable.

The foregoing deduction is based on use of a simple model.

10   According to the detailed first principle calculation by

use of the generalized gradient approximation based on the

density functional method, the formation energy of the In-In

bond is 0.6eV lower than the energy for the In atom in an isolated

state. Energy of a configuration for the two In atoms in the

15   substitutional sites also increases along with distance between

the two In atoms. In other words, the attraction force tends

to provide an effect between the two In atoms. In addition,

the In-In bond is electrically inactive and the empty deep level

deduced by the above-discussed model of the tight-binding

20   approximation appears in the energy gap. On the contrary, for

the B atom, the covalent bond radius is smaller than the Si atom.

In the case of B, the energy of the B-B bond in the nearest neighbor

substitutional site is 0.6 eV higher than the energy for the

B atom in an isolated state. In other words, a repulsive force

25   provided an effect between the two B atoms in the nearest neighbor

substitutional sites. Furthermore, the B-B bond of the nearest

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neighbor substitutional sites is electrically active.

The energy gain $\Delta\varepsilon$ by sp^2 configuring is small in the case of an element such as B having a high electrical activation rate as a dopant. Therefore, the B atoms tend to move away from each other due to the increase of the elastic potential energy. In addition, when the two B atoms are disposed in the nearest neighbor substitutional sites, the B atoms are still electrically active. On the other hand, in the case of a dopant such as In where the energy gain $\Delta\varepsilon$ by sp^2 configuring is large, it is easy for the dopant atoms to come close to each other. In addition, the dopant atoms residing in the nearest neighbor substitutional site are electrically inactive. In other words, In is a dopant which easy to cohere and to be inactive.

Here, the solid solubility limit is considered. The solid solubility limit is defined as the concentration where impurity atoms in the crystal cannot reside in substitutional sites of the crystal lattice so as to cohere and precipitate. Because the precipitation at the beginning concentration of cohesion cannot be observed in an electron microscopes, the solid solubility limit is determined at the upper limit of the carrier concentration in the case of an electrically active dopant. However, in the case of an impurity atom subjected to the sp^2 configuring mechanism, even though the carrier concentration reaches and saturates at the upper limit, the impurity atoms are mainly in the substitutional sites without cohering.

It is conceivable that the combination of the In-In bond

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is replaced in other combination with impurity atoms of In and another kind in order to promote activatation of a dopant in an inactive state. A case where In as the first impurity atom 61 and B as the second, different impurity atom 62 shown in Fig.

5   4, is described using Fig. 6A and 6B.

As shown in Fig. 6A, the case where the first and the second impurity atoms 61, 62 provide the sp^3 configuration in the nearest neighbor substitutional site is considered. Here, the energy level EI_{sp3} of the sp^3 hybridized orbital of the second impurity

10   atom 62 is lower than the energy level EI_{sp3} of the sp^3 hybridized orbital of the first impurity atom 61, and slightly higher than the energy level ES_{sp3} of the sp^3 hybridized orbital of the semiconductor atom 51. The first impurity atom 61 and the semiconductor atom 51 produce the acceptor level E_{A3}. The second

15   impurity atom 62 and the semiconductor atom 51 produce the acceptor level E_{A4}. In addition, the first and the second impurity atoms 61, 62 produce the acceptor level E_{A5}. The acceptor level E_{A4} is slightly shallow in comparison with the acceptor level E_{A3}. In addition, the acceptor level E_{A5} produced by the first and

20   the second impurity atoms is deeper than the acceptor levels E_{A3} and E_{A4}. In other words, in order to be suitable in terms of energy for sp^3 configuring when the second impurity atom 62 is arranged in the nearest neighbor substitutional site of the first impurity atoms 61, the acceptor level E_{A4} which is the energy

25   level of the electronic state produced by the second impurity atom 62 in the substitutional site of the semiconductor crystal,

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is shallower than the acceptor level E_{A3} of the first impurity atoms 61.

On the other hand, as shown in Fig. 6B, when the first and the second impurity atoms 61, 62 provide the sp^2 configuration in the nearest neighbor substitutional site, the energy levels EI_{sp2} of the sp^2 hybridized orbital are low, and the energy level EI_p of the p orbital is high in comparison with the energy level EI_{sp3} of the sp^3 hybridized orbital. Here the energy levels EI_{sp2} and EI_p for the first impurity atom 61 are high in comparison with the second impurity atom 62. Then, acceptor levels E_{A6} and E_{A7} introduced by the combination of the semiconductor atom 51 and the first or the second impurity atoms 61, 62 are produced in the valence band. An acceptor level E_{D3} introduced by the combination of the first and the second impurity atoms 61, 62 is inactive and produced in a deep level near the middle of the band gap.

The electron configuration of the impurity atoms becomes either the sp^3 configuration or the sp^2 configuration depending on magnitude relation between the energy gain $\Delta\varepsilon$ by the sp^2 configuring and the elastic potential energy gain ΔEs. Therefore, it is necessary to make the sp^3 configuration which provides the electrically active acceptor level advantageous in terms of energy by reducing the energy gain $\Delta\varepsilon$ provided by the sp^2 configuring.

Impurity atoms corresponding to such a condition are B and carbon(C) for In in a Si crystal. The covalent bond radii

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of In, B and C are 0.144 nm, 0.088 nm and 0.077 nm, respectively, with relation to Si radius of 0.117 nm. A combination of In and B or In and C provides stress relaxation of the elastic potential energy. In addition, the acceptor levels of In and B are 155

5   meV and 45 meV, respectively, and B has a shallow acceptor level in comparison with In.

Furthermore, C which is a congener element of Si, does not produce the donor or acceptor levels in the substitutional site in the Si crystal. Thus, the energy level of C can be assumed

10   as 0 or a minus value, and satisfies the condition in which the sp^3 configuration becomes advantageous. Actually, according to the result of the first principle calculation, total energies of an In-B cluster and an In-C cluster are 0.6 eV and 0.8 eV, respectively, which are lower than an isolated state of In, B

15   and In, C.

In addition, the In-B bonding and the In-C bonding orbitals are electrically activated. In the case of B, when two B atoms in the substitutional sites and the inactive In-In clusters reside, it is suggested that two active In-B clusters may be

20   formed so as to be stable for approximately 0.6 eV in terms of energy. By the above-mentioned fact, the In-B cluster improves the electrical activation rate. Additionally because of a decrease of energy by the cluster formation, diffusion of the impurity atoms may be suppressed.

25   In other words, by doping In and B with a ratio of 1:1 in a concentration region over the solid solubility limit of

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In, improvement of the activation rate and control of the diffusion profile are possible. This is due to that, in case of the In-C cluster, C does not introduce an acceptor level, and a carrier concentration is half in comparison with B.

5   However, In-C may provide a similar effect as that of the In-B cluster.

The above explanation is an example of a column IV element semiconductor, but the dopant control based on a similar mechanism for a column III-V or a column II-VI compound

10   semiconductor may be possible. In the case of a compound semiconductor, it is necessary to take into account the substitutional sites substituted by a dopant. For example, in the column III-V compound semiconductor, when a group II atom substitutes for a lattice site location of the group III atom,

15   or a group IV atom substitutes for the lattice site location of the group V atom, both the group II atom and group IV atom work as acceptors. In addition, when the group IV atom substitutes for the group III lattice site location, or the group VI atom substitutes for the group V lattice site location, both

20   the group IV atom and group VI atom work as donors. Similarly in the column II-VI compound semiconductor, dopants of plural groups can be used as the acceptors or the donors. In addition, in the column II-VI compound semiconductor, it is substantially different from the column III-V compound semiconductor in that

25   a vacancy has to consider as a dopant. The vacancy works as the acceptor when introduced in the group II lattice site location,

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and as the donor when introduced in the group VI lattice site location.

Next, a description is made of a method to dope the In and B atoms as the first and the second impurity atoms 61, 62 in a semiconductor substrate 10 such as an Si substrate, with respect to the semiconductor device shown in Fig. 1.

At first, on a surface of the semiconductor substrate 10, an insulating film 14 is formed with a thickness of 5 nm by a thermal oxidation method. Through the insulating film 14, ions of In are implanted. Ion implantation conditions are an acceleration energy of 50 keV, and a dose of $1.5*10^{13}cm^{-2}$. Ions of B is implanted afterwards. Ion implantation conditions are an acceleration energy of 7 keV, and a dose of $4*10^{13}cm^{-2}$. Then, a heat treatment is performed after the ion implantations.

In this way, a semiconductor layer 12 having the electrically activated implanted impurity atoms is formed in the surface of the semiconductor substrate 10.

The ions have been implanted, as shown in Fig. 7, so as to have approximately the same peak density of the In and B impurity atoms in the semiconductor substrate 10. The solid solubility limit of In is $1.5*10^{18}cm^{-3}$ at 800-1100 °C. However, by interacting with B, the active In concentration of approximately $6*10^{18}cm^{-3}$ corresponding to the peak ion-implanted concentration of B can be achieved. Therefore, a total amount of the active impurity concentration near the region of the peak ion-implanted concentration of approximately $1*10^{19}cm^{-3}$ can be achieved.

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Therefore, effect of the activation rate by interaction of In

and B applied to a channel region of a MOSFET can be improved.

For example, as shown in Fig. 7, the channel region having

the steep concentration distribution so as to have a low impurity

5   concentration of approximately $2*10^{18}$ cm^{-3} at the surface portion

of the semiconductor substrate 10, and to have an active impurity

concentration of more than $1*10^{19}$ cm^{-3} at approximately 20 nm

deep from the surface.

In the embodiment of the present invention, the ion

10   implantation energy of In and B are determined where the peak

of the active impurity concentrations are located at

approximately 20 nm deep from the surface of the semiconductor

substrate 10.   However, the peak location of the activity

impurities concentration can be provided optionally by

15   appropriately setting the ion implantation energy.   In addition,

in the embodiment of the present invention, the ion implantation

peak concentrations of In and B are set at the same level.   However,

because it is an essential attribute that the B concentration

corresponding to the desired activity impurities concentration

20   is included in a region increasing the activity of the In

concentration, the ion implantation peak position of each can

be set optionally.   In addition, the peak concentration can be

set to the optional concentration by adjusting the dose of the

impurity atom.

25   According to the embodiment of the present invention, the

In and B impurity atoms may become electrically active by arranging

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In and B in the nearest neighbor substitutional site in the Si crystal. Furthermore, for decrease of energy by clustering between In and B, generation of the In-B cluster is enhanced, and the diffusion of In and B is suppressed.

5   Next a method to dope In and B in a source/drain region of a p-channel MOSFET is described by use of Figs. 8A through 8F.

(a)   As shown in Fig. 8A, an oxide film 24 of 5 nm thick is formed by a thermal oxidation method on a surface of a
10   semiconductor substrate 20 of an n type Si. Then, an impurity doping layer 22 doped with donor impurity atoms such as As and P as a 20 nm deep channel region is formed by ion implantation.

(b)   On the oxide film 24, as shown in Fig. 8B, a conductive poly-silicon film 26 is deposited.

15   (c)   By using a photolithography technique, as shown in Fig. 8C, the poly-silicon film 26 is processed to form a gate electrode 36. Then the oxide film 24 beneath the gate electrode 36 is formed as a gate oxide film 34 with the same thickness of 5 nm. For the region apart from the gate electrode 36, the
20   oxide film 24 is formed as a thinner oxide film 24a by a gate electrode formation process.

(d)   In the semiconductor substrate 20, each of the impurity atoms of In and B is ion-implanted sequentially with peak impurity concentrations of approximately $1*10^{19}$ cm^{-3}, and
25   peak positions of approximately 30 nm deep. By an annealing process, as shown in Fig. 8D, an extension diffusion layer 42

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is formed. Here, In and B are not implanted into the impurity

doping layer 22 beneath the gate oxide film 34 so that the gate

electrode 36 works as a mask of the ion implantation.

(e) Subsequently, a thick oxide film is deposited by a

5   chemical vapor deposition (CVD) method or the like. By

directional etching, such as reactive ion etching, as shown in

Fig. 8E, a side wall insulating film 38 is formed in a side wall

of the gate electrode 36.

(f) Then, In and B are ion-implanted using the gate

10 electrode 36 and the side wall insulating film 38 as a mask.

By an annealing process, as shown in Fig. 8F, a source/drain

diffusion layer 44 is formed. Each of the impurity atoms of

In and B is ion-implanted sequentially with peak impurity

concentrations of approximately $1*10^{20}$ cm^{-3}, and peak positions

15 of approximately 100 nm deep. The impurity doping layer 22

beneath the gate electrode 36 and the extension diffusion layer

42 beneath the side wall insulating film 38 are masked by the

gate electrode 36 and the side wall insulating film 38, so as

not to be affected by formation of the source/drain diffusion

20 layer 44.

As mentioned above, by the interaction of In and B, the

active concentration of In increases. Furthermore, since the

diffusion of In and B is suppressed, the extension diffusion

layer 42 and the source/drain diffusion layer 44 may be formed

25 in a desired region with a desired active impurity concentration.

(OTHER EMBODIMENTS)

The present invention has been described as mentioned above. However the descriptions and drawings that constitute a portion of this disclosure should not be perceived as limiting this invention.

5 Various alternative embodiments and operational techniques will become clear to persons skilled in the art from this disclosure.

For example, after doping In and B in a Si substrate using ion implantation and the like, a Si epitaxial growth layer may be grown on the Si substrate surface so as to form the SSRP which

10 has a lower surface impurity concentration. Additionally, it is permissible that, before the epitaxial growth on a Si substrate, either of In or B is ion-implanted first, and after the epitaxial growth of the Si layer, the other is ion-implanted. In addition, after doping either In or B, or both in a Si substrate, a portion

15 of the Si substrate is etched and then the epitaxial growth of the Si layer is conducted so as to form the SSRP having a great impurity concentration difference.

In addition, in the embodiment of the present invention, the Si crystal doped with In and B has been described. However,

20 any atom having a small or large covalent bond radius compared with the covalent bond radius of Si, is at least a dopant for Si, and the dopant atom and other atom are inactive in the nearest neighbor lattice site location. Then, the same effect as the embodiment of the present invention is provided. Furthermore,

25 instead of Si, for example, for a column IV semiconductor such as Ge, a SiGe system, a SiGeC system, a column III-V compound

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semiconductor such as gallium arsenide (GaAs), aluminum nitride (AlN), gallium nitride (GaN), indium gallium nitride ($In_xGa_{1-x}N$), a column II-VI compound semiconductor such as zinc oxide (ZnO), zinc sulfide (ZnS), and the like, when dopant atoms have the same

5   magnitude relation as Si with a covalent bond radius of a constituent atom of the semiconductor and are also inactive in the nearest neighbor substitutional site, then the same effect as the embodiment of the present invention is provided. In particular, in a diamond crystal, a combination of P as the first impurity atom and nitrogen

10   (N) as the second impurity atom is desirable. In addition, in a GaN crystal, a combination of beryllium (Be) as the first impurity atom and C as the second impurity atom is desirable.